

# 'The Gc Ms Analysis Of Ethyl Acetate Extract Of One Herbal Plant, 'Justicia Glauca'

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#### ABSTRACT

The present study deals with the GC MS analysis of one medicinal plant, Justiciaglauca. Not much work was done on this species although all the species of Justicia genus have important ethno-medical use. The plant was collected from nearby hills of Chengalpattu, Tamilnadu. The ethyl acetate extract of the aerial parts of the plant was subjected to GC MS study following standard protocols. It was observed that some very important molecules such asn-Hexadecanoic acid, 3,7,11,15-Tetramethyl-2-hexadecen-1-ol, 2-((Octan-2-yloxy)carbonyl)benzoic acid, Squalene, Sulfurous acid, butyl heptadecyl ester, .gamma.-Tocopherol, Pregnenolone, 17.alpha.-Hydroxypregnenolone, Stigmasterol, .beta.-Sitosterol, Z,E-2,13-Octadecadien-1-ol, .beta.-Amyrin, Lupeol. Further work is warranted in this regard. These molecules play a vital role in the medicinal role of Justiciaglauca.

Key Words GC MS, Justiciaglauca, n-Hexadecanoic acid, Stigmasterol, .beta.-Sitosterol,.beta.-Amyrin, Lupeol

## INTRODUCTION

Justicia genus has a number Of species which have great medicinal roles. Justiciaadhatoda is one plant which has maximum ethno-medicinal value. Only scanty reports are available on the medicinal

roles of other related species of this genus. The phytochemical and analysis and antibacterial activity of Justiciaglaucawas reported by Bheemaganiet al, 2015. The antibacterial role of the gold nanoparticles of Justiciaglauca was studied by Emmanuel et al, 2017. Kavithaet al, 2014 have reviewed the phytochemical and pharmacological potential of a related species, Justiciagendarussa. The present work reports the GC MS pattern of the ethyl acetate extracts of Justiciagaluca whole plant. This is in continuation of our endeavour to establish the medicinal efficacy of the herbal and traditional systems of Ayurveda, Sidhha and Unani systems of medicine (Priyadarshiniet al, 2017; Jayakumariet al, 2017; Raoet al, 2018; Vijayalakshmi and Rao, 2019; Yuvarajet al, 2019; Mutteviet al, 2019, Raoet al, 2019; Mutteviet al, 2020; Vijayalakshmi and Rao, 2020; Janakiet al, 2021, Perumalet al, 2021).

## MATERIALS AND METHODS

The plant Justiciaglaucawas collected from the nearby hills at Chengalpattu, Tamil Nadu. The plant was identified by a qualified botanist at Chennai. The ethyl acetate extract of the shade dried whole plant was collected after 48 h of soaking. The extract was evaporated and the dried powder was used for GC-MS analysis by standard procedures.

## GC-MS Procedure

Instrument: GC (Agilent: GC: (G3440A) 7890A. MS/MS: 7000 Triple Quad GCMS) was equipped with MS detector.

## Sample Preparation

About 100 ml sample was dissolved in 1 ml of suitable solvents. The solution was stirred vigorously using vortex stirrer for 10 s. The clear extract was determined using GC for analysis.

#### **GC-MS Protocol**

Column DB5 MS (30 mm × 0.25 mm ID ×0.25  $\mu$ m, composed of 5% phenyl 95% methylpolysiloxane), electron impact mode at 70 eV; helium (99.999%) was used as carrier gas at a constant flow of 1 ml/min injector temperature 280°C; auxilary temperature: 290°C ion-source temperature 280°C.

The oven temperature was programmed from 50°C (isothermal for 1.0 min), with an increase of 40°C/min, to 170°C C (isothermal for 4.0 min), then 10°C/min to 310°C (isothermal for 10 min) fragments from 45 to 450 Da. Total GC running time is 32.02 min. The compounds are identified by GC-MS Library (NIST and WILEY).

# **RESULTS AND DISCUSSION**

The results of the GC-MS analysis of the whole plant ethyl acetate extract, along with the possible medicinal role of each molecule of Justiciaglaucaextract are tabulated in Table 1. Figure 1 represents the GC-MS profile of ethyl acetate extract of the whole plant of Justiciaglauca. The identification of metabolites was accomplished by comparison of retention time and fragmentation pattern with mass spectra in the NIST spectral library stored in the computer software (version 1.10 beta, Shimadzu) of the GC-MS along with the possible pharmaceutical roles of each bio molecule as per Dr. Duke's Phytochemical and ethno-botanical data base (National Agriculture Library, USA) and others as shown in Table 1.Table no. 1 indictes the presence of a wide range of molecules having important medicinal roles, such as n-Hexadecanoic acid, 3,7,11,15-Tetramethyl-2-hexadecen-1-ol, 2-((Octan-2-yloxy)carbonyl)benzoic acid, Squalene, Sulfurous acid, butyl heptadecyl ester, .gamma.-Tocopherol, Pregnenolone, 17.alpha.-Hydroxypregnenolone, Stigmasterol, .beta.-Sitosterol, Z,E-2,13-Octadecadien-1-ol, .beta.-Amyrin, Lupeol. Further work is warranted in this regard.

### CONCLUSION

It is concluded that the types of medicinal values the molecules have, as is shown in the Table, Justiciagaluca can prove itself as an important medicinal plant.

## ACKNOWLEDGMENTS

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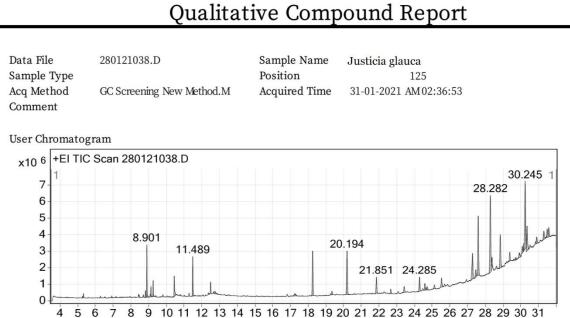
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Figure 1. Represents the GC MS graph of ethyl acetate extract Justiciaglauca'.



10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 Counts vs. Acquisition Time (min) 6 8 9 5 7

Table 1. Indicates the retentions time, types of possible compound, molecular formula, molecular mass, percentage peak area and the possible medicinal roles of each compound as shown in the GC MS profile ofJusticiaglauca'.

Ret.	Compound	Mol.	Mol.	% Peak	Possible Medicinal Role
Time		Formula	Mass	Area	
8.90	Bicyclo[3.1.1]heptane,	C10H18	138.1	4.18	Not Known
	2,6,6-trimethyl-				
9.25	9-Octadecyne	C18H34	250.3	1.27	Not Known
10.45	n-Hexadecanoic acid	C16H32O	256.2	2.57	Acidifier, Arachidonic acid
		2			Inhibitor, Increases Aromatic
					Amino acid decarboxylase
					activity, Inhibits production
					of uric acid, Urine acidifier,

					Anaphylactic, Arylamine N acetyltransferase inhibitor, decreases norepinephrine production, Down regulates nuclear and cytosol androgen reuptake, GABA-nergic, Increase NK cell activity, inhibits production of tumor necrosis factor, Myo-neuro- stimulator
11.49	Cyclohexan ol, 5- methyl-2- (1- methylethy l)-, (1.alpha.,2. beta.,5.alp ha.)-(.+/)-	C10H20O	156.2	3.98	Not Known
12.50	3,7,11,15-Tetramethyl-2- hexadecen-1-ol	C20H40O	296.3	1.29	Oligosaccharide Provider
18.26	2-((Octan-2- yloxy)carbonyl)benzoic acid	C16H22O 4	278.2	6.61	Acidifier, Arachidonic acid inhibitor, Increases Aromatic Amino acid Decarboxylase activity
20.19	Squalene	C30H50	410.4	7.01	Monooxygenase inhibitor, biochemical precursor in the preparation of steroids, natural moisturizer, used in cosmetics
21.85	Sulfurous acid, butyl heptadecyl ester	C21H44O 3S	376.3	2.69	Acidifier, Arachidonic acid inhibitor, Increases Aromatic Amino acid Decarboxylase

					activity
23.42	1-Nonylcycloheptane	C16H32	224.3	1.13	Not known
24.59	Octacosyl acetate	C30H60O 2	452.5	1.17	Not known
24.71	.gammaTocopherol	C28H48O 2	416.4	0.71	Tocopherol synergist, PPAR-gamma antagonist
25.53	4,5,6,7-Tetrahydro- benzo[c]thiophene-1- carboxylic acid allylamide	C12H15N OS	221.1	1.64	Not known
27.27	Pregnenolone	C21H32O 2	316.2	5.83	Helps relieve fatigue and provides energy, enhances memory; trauma and improves immunity, used for skin disorders including psoriasis and scleroderma.
27.46	17.alpha Hydroxypregnenolone	C21H32O 3	332.2	1.44	5 alpha reductase inhibitor, HIF1 alpha inhibitor, Alpha amylase inhibitor, IkappaB- alpha phosphorylation inhibitor, Interlukine- 1 alpha inhibitor, Testosterone 5 alpha reductase inhibitor, alpha agonist, alpha amylase inhibitor, alpha glucoside inhibitor, increases alpha Mannosidase activity, TNF alpha inhibitor
27.59	Stigmasterol	С29Н48О	412.4	11.90	Precursor of progesterone , acts as intermediate in the biosynthesis of androgens and estrogens, anti- osteoarthritic,

28.28.betaSitosterolC29H500414.414.5317 beta dehydrogenase inhibitor, androgen blocker, anti-androgen blocker, anti-anti-angyloid beta, anti-angyloid beta, Beta 2- receptor, beta blocker, beta-galactosidase inhibitor28.37Z,E-2,13-Octadecadien-1-olC18H340266.31.30Increases Zinc bioaxilability, provides zinc, antitamyci or vides zinc, antitamor, Cytochrome-P450-2E1- Inhibitor, Decreases C- Teleopeptide Excretion, Decreases C-<						antihypercholesterolemic,
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28.28.betaSitosterolC29H500414.414.5317 beta dehydrogenase inhibitor, androgen blocker, anti-amyloid beta, anticancer, Anti TGF beta, Beta 2- receptor, beta blocker, beta-galactosidase inhibitor28.37Z,E-2,13-Octadecadien-1-olC18H340266.31.30Increases Zinc bioavailability, provides zinc, anticancer, antidote, antitumor, Cytochrome-P450-2E1- Inhibitor, Decreases C- Teleopeptide Excretion, Decreases Deoxypyrdinoline Excretion, Decreases Epinephrine Production, Decreases Oxalate Excretion, Decreases Oxalate, Anti TGF Deta, Beta adrenergic receptor Blocker, beta blocker, beta blocker, beta blocker, beta						hypoglycemic, antimutagenic,
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29.38.betaAmyrinC30H500426.41.3817 beta hydroxysteroid dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta adrenergic receptor blocker, beta blocker, beta						provides zinc, anticancer,
29.38.betaAmyrinC30H500426.41.3817 beta hydroxysteroid deta, Anti TGF beta, Beta adrenergic receptor blocker, beta blocker, beta						antidote, antitumor,
29.38.betaAmyrinC30H500426.41.3817 beta hydroxysteroid dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta, Beta adrenergic receptor blocker, beta blocker, beta						Cytochrome-P450-2E1-
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29.38.betaAmyrinC30H500426.41.3817 beta hydroxysteroid dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta, Beta adrenergic receptor blocker, beta blocker, beta						Teleopeptide Excretion,
29.38.betaAmyrinC30H500426.41.3817 beta hydroxysteroid dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta, Beta adrenergic receptor blocker, beta blocker, beta						Decreases Deoxypyridinoline
29.38.betaAmyrinC30H500426.41.3817 beta hydroxysteroid dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta, Beta adrenergic receptor blocker, beta blocker, beta						Excretion, Decreases
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29.38.betaAmyrinC30H500426.41.3817 beta hydroxysteroid dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta, Beta receptor agonist, Beta adrenergic receptor blocker, beta blocker, beta						Adhesion, Decreases
29.38.betaAmyrinC30H500426.41.3817 beta hydroxysteroid dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta, Beta receptor agonist, Beta adrenergic receptor blocker, beta blocker, beta						Epinephrine Production,
dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta, Beta receptor agonist, Beta adrenergic receptor blocker, beta blocker, beta						Decreases Oxalate Excretion
dehydrogenase inhibitor, Antiamyloid beta, Anti TGF beta, Beta receptor agonist, Beta adrenergic receptor blocker, beta blocker, beta						
Antiamyloid beta, Anti TGF beta, Beta receptor agonist, Beta adrenergic receptor blocker, beta blocker, beta	29.38	.betaAmyrin	C30H50O	426.4	1.38	17 beta hydroxysteroid
beta, Beta receptor agonist, Beta adrenergic receptor blocker, beta blocker, beta						dehydrogenase inhibitor,
Beta adrenergic receptor blocker, beta blocker, beta						Antiamyloid beta, Anti TGF
blocker, beta blocker, beta						beta, Beta receptor agonist,
						Beta adrenergic receptor
galactosidase inhibitor, beta						blocker, beta blocker, beta
						galactosidase inhibitor, beta

					glucuronidase inhibitor, ER
					beta binder
30.08	2-[4-methyl-6-(2,6,6-	C23H32O	324.2	1.06	NotKnown
	trimethylcyclohex-1-				
	enyl)hexa-1,3,5-				
	trienyl]cyclohex-1-en-				
	1-carboxaldehyde				
30.25	Lupeol	C30H50O	426.4	12.37	Anti-inflammatory, anti-
					arthritic, anti-mutagenic and
					anti-malarial